

Surgical Placement of BioTend During an Open Surgery

1. Selection of BioTend size and placement at repair site

The surgical preparation of rotator cuff repair, including placement of repair sutures through tendon is completed, but attachment to bone anchors is not done, therefore allowing placement of sutures for BioTend.

2. Select width of BioTend

BioTend should cover the width of the rotator cuff tear repair, therefore providing mechanical augmentation across all of the tendon-to-bone repair line, and also any tendon-to-tendon repair (for example, margin convergence).

3. Identify location of medial tendon sutures

These should be positioned medially to repair sutures, laterally to muscle, and within the projected location of BioTend.

4. Place sutures in tendon

The selection of number and type of suture is at the surgeon's discretion, and some recommendations are below. More sutures placed medially and laterally will enhance the functionality of the device.

- 12 mm device: 2 simple sutures separated by 8 mm
- 25 mm device: 3 - 4 simple sutures at 5 mm intervals
- 25 mm device: 2 simple sutures placed 5 mm from edge, 1 mattress suture centrally
- 25 mm device: 3 mattress sutures, placed 5 mm from edge and centrally.
- 40 mm device, 4 – 6 simple sutures
- 40 mm device: 2 simple, 1 mattress stitch, 2 simple

When these sutures are in place, the rotator cuff repair should be completed.

5. Placement of lateral bone anchors

The objective is to maximize uniform attachment of BioTend to the bone. Bone anchors (preferably threaded to maximize load-bearing potential) are usually used to fix BioTend laterally to the bone. These may be anchors shared with fixing the rotator cuff tendon to the bone, or may be anchors dedicated to attaching BioTend to bone. The anchors should be placed laterally to the rotator cuff repair (if dedicated to BioTend), or at the lateral position of the repair (if shared with rotator cuff repair). The anchors should be positioned evenly over the projected position of BioTend, to uniformly distribute the load between the device and the anchors. There is flexibility in how many sutures, and what type of stitch is used. For example:

- 12 mm device: 1 bone anchor to support 2 simple sutures
- 25 mm device: 2 bone anchors to support 4 simple sutures
- 25 mm device: 2 bone anchors to support 4 mattress sutures
- 25 mm device: 2 bone anchors to support 2 mattress sutures
- 25 mm device: 3 bone anchors to support 3 mattress sutures
- 25 mm device: 3 bone anchors to support 6 simple sutures
- 25 mm device: 3 bone anchors to support 1 mattress suture centrally, 4 simple sutures anterior and posterior
- 40 mm device: 3 bone anchors to support 6 simple sutures
- 40 mm device: 3 bone anchors to support 3 - 6 mattress sutures

6. **Select length of BioTend**

The objective is to select the length of BioTend so that it extends from the medial sutures to the lateral bone anchors.

- A bite size of >5 mm is used to place the sutures internal to the welded ends of BioTend.
- The type of suture selected, particularly at the bone anchor site, provides flexibility in selection of the length of BioTend.

For example, if BioTend extends over the bone anchors then mattress stitches are more likely to be used, and if BioTend does not cover the bone anchors, either mattress, simple or a combination of both may be used. Simple sutures overlapping the end of BioTend will prevent any tendency of the lateral end of the device to lift from the bone surface.

7. **Suture BioTend to anchors and tensioning**

The lateral sutured are completed, ensuring tension is applied along the length of BioTend, to ensure load sharing over the rotator cuff repair site. Additional sutures along the anterior and posterior edges of BioTend can then be placed for additional fixation strength.